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NEWS		JUN		USPATFULL and USPAT2 updated with 11-character
	•	0011		patent numbers for U.S. applications
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NEWS	9	JUN	30	STN on the Web enhanced with new STN AnaVist Assistant and BLAST plug-in
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NEWS		JUL		CA/CAplus patent coverage enhanced
NEWS		JUL		EPFULL enhanced with additional legal status
				information from the epoline Register
NEWS	13	JUL	28	IFICDB, IFIPAT, and IFIUDB reloaded with enhancements
NEWS				STN Viewer performance improved
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NEWS		AUG		CA/CAplus enhanced with printed Chemical Abstracts
		1100		page images from 1967-1998
NEWS	17	AUG	1.5	CAOLD to be discontinued on December 31, 2008
NEWS		AUG		CAplus currency for Korean patents enhanced
NEWS		AUG		CAS definition of basic patents expanded to ensure
CMEN	19	AUG	21	cas definition of basic patents expanded to ensure comprehensive access to substance and sequence information
NEWS	20	SEP	18	Support for STN Express, Versions 6.01 and earlier, to be discontinued
NEWS	21	SEP	25	CA/CAplus current-awareness alert options enhanced to accommodate supplemental CAS indexing of exemplified prophetic substances
NEWS	22	SEP	26	WPIDS, WPINDEX, and WPIX coverage of Chinese and and Korean patents enhanced
NEWS	23	SEP	29	IFICLS enhanced with new super search field
NEWS	2.4	SEP		EMBASE and EMBAL enhanced with new search and
				display fields
NEWS	25	SEP	30	CAS patent coverage enhanced to include exemplified prophetic substances identified in new Japanese-language patents
NEWS	26	OCT	0.7	EPFULL enhanced with full implementation of EPC2000
NEWS		OCT		Multiple databases enhanced for more flexible patent number searching

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=> relaxin-3 L1 199 RELAXIN-3

=> 11 and feed?

19 L1 AND FEED? => 11 and weight

=> 12 or 13 22 L2 OR L3 L4

L3

=> dup rem 14 PROCESSING COMPLETED FOR L4

12 DUP REM L4 (10 DUPLICATES REMOVED)

=> 15 and 1970-2005/pv 4 L5 AND 1970-2005/PY L6

=> d ibib 16 1-4

L6 ANSWER 1 OF 4 MEDLINE on STN ACCESSION NUMBER: 2005355604 MEDI-THE DOCUMENT NUMBER: PubMed ID: 15845619

8 L1 AND WEIGHT

TITLE: Central relaxin-3 administration causes hyperphagia in male Wistar rats.

AUTHOR: McGowan B M C; Stanley S A; Smith K L; White N E; Connolly

M M; Thompson E L; Gardiner J V; Murphy K G; Ghatei M A;

Bloom S R

CORPORATE SOURCE: Endocrine Unit, Imperial College School of Medicine, Hammersmith Hospital, London W12 ONN, United Kingdom.

SOURCE: Endocrinology, (2005 Aug) Vol. 146, No. 8, pp.

3295-300. Electronic Publication: 2005-04-21.

Journal code: 0375040. ISSN: 0013-7227.

PUB. COUNTRY: United States

DOCUMENT TYPE: Journal; Article; (JOURNAL ARTICLE) (RESEARCH SUPPORT, NON-U.S. GOV'T)

LANGUAGE: English

FILE SEGMENT: Abridged Index Medicus Journals; Priority Journals

ENTRY MONTH: 200509

ENTRY DATE: Entered STN: 13 Jul 2005

Last Updated on STN: 27 Sep 2005

Entered Medline: 26 Sep 2005

L6 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2005:1289617 CAPLUS

DOCUMENT NUMBER: 144:32547

TITLE: Methods and compositions for control of fetal growth

via modulation of relaxin Unemori, Elaine
PATENT ASSIGNEE(S):
Bas Medical, Inc., USA

SOURCE: PCT Int. Appl., 93 pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

	PATENT NO.																	
WO	2005	1154	35					1					20050502 <					
	W:	CN, GE,	CO, GH,	CR, GM,	CU, HR,	CZ, HU,	DE, ID,	AZ, DK, IL,	DM, IN,	DZ, IS,	EC, JP,	EE, KE,	EG, KG,	ES, KM,	FI, KP,	GB, KR,	GD, KZ,	
		NI,	NO, SY,	NZ,	OM,	PG,	PH,	LV, PL, TT,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	
	RW:	BW, AZ, EE,	GH, BY, ES,	KG, FI,	KZ, FR,	MD, GB,	RU, GR,	MZ, TJ, HU, BJ,	TM, IE,	AT, IS,	BE, IT,	BG, LT,	CH, LU,	CY, MC,	CZ,	DE, PL,	DK, PT,	
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CA US	2563 2006	433 0247	163		A1 A1		2005 2006	1208 1102	CA 2005-2563433 US 2005-120582 EP 2005-780049					20050502				-
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US US	JP 2007535574 US 20060247172 US 20080108572 PRIORITY APPLN. INFO::						2006	1102	1	US 2 US 2	007- 006- 007- 004-	4782 9813	67 38		2	0060	628 031	

US 2005-120582 A3 20050502

W 20050502

WO 2005-US15248

L6 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2005:1016407 CAPLUS

DOCUMENT NUMBER: 144:101114

TITLE: The relaxin gene-knockout mouse: a model of

progressive fibrosis

AUTHOR(S): Samuel, Chrishan S.; Zhao, Chongxin; Bathgate, Ross A. D.; Du, Xiao-Jun; Summers, Roger J.; Amento, Edward P.; Walker, Leslev L.; McBurnie, Mary; Zhao, Ling;

Tregear, Geoffrey W.

CORPORATE SOURCE: Howard Florey Institute of Experimental Physiology &

Medicine, University of Melbourne, Parkville, Victoria, 3010, Australia

SOURCE: Annals of the New York Academy of Sciences (

2005), 1041 (Relaxin and Related Peptides),

173-181

CODEN: ANYAA9; ISSN: 0077-8923 PUBLISHER: New York Academy of Sciences DOCUMENT TYPE: Journal; General Review

LANGUAGE: English

REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2008 ACS on STN 2005:823818 CAPLUS

ACCESSION NUMBER: DOCUMENT NUMBER: 143:223700

TITLE: Binding of relaxin-3 to

G-protein-coupled receptor 135 (GPCR135) and a role in promoting food intake and body weight gain and

obesity: use for antiobesity drug screening

INVENTOR(S): Hida, Takayuki; Hirohashi, Tomoko; Sawai, Toru; Seiki, Takashi; Takahashi, Eiki; Kosasa, Michiko; Harada,

Kokichi; Arai, Toru

PATENT ASSIGNEE(S): Eisai Co., Ltd., Japan SOURCE: PCT Int. Appl., 97 pp. CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1 PATENT INFORMATION:

PATENT NO.				KIND DATE			APPLICATION NO.											
WO 2005075641					A1		2005	0818										
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ΑU	2005	2103	69		A1		2005	0818		AU 2	005-	2103	69		2	0050	209 <	
CA	2555	469			A1		2005	0818		CA 2	005-	2555	469		2	0050	209 <	
EP	1721	971			A1		2006	1115		EP 2	005-	7099	39		2	0050	209	
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		HR,	MK,	YU														

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CN 1918290 A 20070221 CN 2005-80004472 20050209
US 20070054850 A1 20070308 US 2006-588542 20060807
KR 2007004699 A 20070109 KR 2006-718442 20060908
KITY APEN. INFO: JP 2004-31591 A 20040209
PRIORITY APPLN. INFO.:
                                                JP 2004-368509 A 20041220
WO 2005-JP1887 W 20050209
REFERENCE COUNT: 11 THERE ARE 11 CITED REFERENCES AVAILABLE FOR THIS
                                  RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
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=> tomoko?/au and sekiva?/au
            0 TOMOKO? AU AND SEKIYA? AU
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=> 112 and 113
            0 L12 AND L13
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L18
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T.19
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     ANSWER 1 OF 4 MEDLINE on STN
L6
     Relaxin-3 (INSL-7) is a recently discovered member of
AB
     the insulin superfamily. Relaxin-3 mRNA is expressed in the nucleus incertus of the brainstem, which has projections to the
     hypothalamus. Relaxin-3 binds with high affinity to
     the LGR7 receptor and to the previously orphan G protein-coupled receptor
     GPCR135. GPCR135 mRNA is expressed predominantly in the central nervous
     system, particularly in the paraventricular nucleus (PVN). The presence
     of relaxin-3 and these receptors in the PVN led us to
     investigate the effect of central administration of relaxin-
     3 on food intake in male Wistar rats. The receptor involved in
     mediating these effects was also investigated. Intracerebroventricular
     injections of human relaxin-3 (H3) to satiated rats
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significantly increased food intake 1 h post administration in the early light phase [0.96 +/- 0.16 g (vehicle) vs. 1.81 +/- 0.21 g (180 pmol H3), P < 0.05] and the early dark phase [2.95 +/- 0.45 g (vehicle) vs. 4.39 +/-0.39 g (180 pmol H3), P < 0.05]. Intra-PVN H3 administration significantly increased 1-h food intake in satiated rats in the early light phase [0.34 +/- 0.16 g (vehicle) vs. 1.23 +/- 0.30 g (18 pmol H3), P < 0.05] and the early dark phase [4.43 +/- 0.32 g (vehicle) vs. 6.57 +/-0.42 g (18 pmol H3), P < 0.05]. Feeding behavior increased after intra-PVN H3. Equimolar doses of human relaxin-2, which binds the LGR7 receptor but not GPCR135, did not increase feeding. Hypothalamic neuropeptide Y, proopiomelanocortin, or agouti-related peptide mRNA expression did not change after acute intracerebroventricular H3. These results suggest a novel role for relaxin-3 in appetite regulation.

- ANSWER 2 OF 4 CAPLUS COPYRIGHT 2008 ACS on STN AB The invention relates to the method for treatment, diagnosis and prevention of diseases related to fetal growth and placental insufficiency and comprises methods including inhibiting or increasing relaxin synthesis, relaxin receptor synthesis, relaxin binding to the relaxin receptor, and relaxin receptor activity. The invention also relates to screening assays to identify compds. that modulate relaxin and/or relaxin receptor activity. The invention further relates to gene therapy methods
 - utilizing relaxin and relaxin-related sequences for the treatment and prevention of diseases related to fetal growth and placental insufficiency.
- ANSWER 3 OF 4 CAPLUS COPYRIGHT 2008 ACS on STN
- AB A review. Relaxin is well known for its actions on collagen remodeling. To improve our understanding of the physiol. role(s) of relaxin, the relaxin gene-knockout (RLX-KO) mouse was established by our group and subsequently phenotyped. Pregnant RLX-KO mice underwent inadequate development of the pubic symphysis as well as the mammary glands and nipples compared to wild-type mice, thus preventing lactation. Later studies showed that these deficiencies were associated with increased collagen, primarily in the nipple and vagina. Anal. of male RLX-KO mice also demonstrated inadequate reproductive tract development. The testis, epididymis, and prostate of RLX-KO mice showed delayed tissue maturation and growth associated with increased collagen deposition. In nonreproductive tissues, an age-related increase in interstitial collagen (fibrosis) was also detected in the lung, heart, and kidneys of RLX-KO mice and was associated with organ dysfunction. From 6-9 mo of age and onwards, all organs of RLX-KO mice, particularly male mice, underwent progressive increases in tissue weight and collagen content (all P <.05) compared with wild-type animals. The increased fibrosis contributed to bronchiole epithelium thickening and alveolar congestion (lung), atrial hypertrophy and increased ventricular chamber stiffness (heart) in addition to glomerulosclerosis (kidney). Treatment of RLX-KO mice with recombinant human relaxin in early and developed stages of fibrosis caused the reversal of collagen deposition in the lung, heart, and kidneys. Together, these findings suggest that relaxin is a naturally occurring inhibitor of collagen deposition during normal development, aging, and pregnancy and can be used to prevent the progression of fibrosis.
- ANSWER 4 OF 4 CAPLUS COPYRIGHT 2008 ACS on STN
 - The present invention provides a polypeptide having useful effects of promoting food intake and body weight gain and causing obesity; drugs for diseases containing this polypeptide; a method of screening substances activating or inhibiting a receptor for the polypeptide; and food intake regulating agents, drugs for obesity, diabetes and so on containing a substance inhibiting the expression of the polypeptide, etc. By intracerebroventricularly administering relaxin-3 to

rats and observing feed intake, body weight, fat amount, etc. after the administration, it was found that relaxin—3 has effects of promoting food intake and body weight gain and cause obesity.

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(FILE 'HOME' ENTERED AT 23:09:28 ON 14 OCT 2008)
FILE 'MEDLINE, BIOSIS, CAPLUS' ENTERED AT 23:09:53 ON 14 OCT 2008 L1 199 RELAXIN-3 L2 19 L1 AND FEED? L3 8 L1 AND WEIGHT L4 22 L2 OR L3 L5 12 DUP REM L4 (10 DUPLICATES REMOVED) L6 4 L5 AND 1970-2005/PY L7 0 TAKAYUKIT/AU AND HIDA?/AU L8 0 TOMOKO?/AU AND SEKIYA?/AU L9 0 TORU?/AU AND SAWAI?/AI L9 0 TORU?/AU AND SAWAI?/AI L10 0 TORU?/AU AND SAWAI?/AI
L11 0 TORO?/AU AND SAWA1?/AU L11 0 TAKASHI?/AU AND SEIKI?/AI
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L13 2 EIKI?/AU AND TAKAHASHI?/AU
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L15 0 KOKICHI?/AU AND HARADA?/AU
L16 0 TOHRU?/AU AND ARAI?/AU
L17 0 L12 AND L13
L18 11 L12 OR L13
L19 0 L6 AND L18
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FULL ESTIMATED COST 68.72 68.93
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS) SINCE FILE TOTAL ENTRY SESSION

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-2.40

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